California Drug Recall Information

Recall Name
Allergan Recalls Ophthalmic Products Due to Particulate Matter

Recall Date	Product Description	Recalling Firm	Recall Reason
8/24/15	• **REFRESH® Lacri-Lube®** 3.5g and 7g ***NDC # 0023-0312-07*** • **REFRESH P.M.®** 3.5g ***NDC # 0023-0240-04*** • **FML®** (fluorometholone ophthalmic ointment) 0.1%, 3.5g ***NDC # 0023-0316-04*** • **Blephamide®** (sulfacetamide sodium and prednisolone acetate ophthalmic ointment, USP) 10% / 0.2%, 3.5g ***NDC # 0023-0313-04***	**Allergan** Dublin, Ireland	*Due to complaints which reported a small black particle, identified as part of the cap, which could potentially be introduced into the product.*

Recall Class	Product Identification	Distribution	Affected Dates
N/A	Click for Affected Lots / Expiration Dates. Product Labels	**CA**, nationwide	Expiration Dates: Apr-17 to Mar-18

FOR ADDITIONAL INFORMATION, PLEASE VISIT:

http://www.fda.gov/Safety/Recalls/ucm459485.htm